

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
21 June 2001 (21.06.2001)

PCT

(10) International Publication Number  
WO 01/43788 A2

(51) International Patent Classification<sup>7</sup>: A61L 27/34, 27/54, 29/08, 29/16, 31/10, 31/16

(21) International Application Number: PCT/US00/42372

(22) International Filing Date:  
29 November 2000 (29.11.2000)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:  
09/461,846 15 December 1999 (15.12.1999) US

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(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SB, SG, SI, SK, SL, TI, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:  
— Without international search report and to be republished upon receipt of that report.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.



WO 01/43788 A2

(54) Title: POLYMER COMPOSITIONS CONTAINING COLLOIDS OF SILVER SALTS

(57) Abstract: The present invention relates to antimicrobial compositions, methods for the production of these compositions, and use of these compositions with medical devices, such as catheters, and implants. The compositions of the present invention advantageously provide varying release kinetics for the active ions in the compositions due to the different water solubilities of the ions, allowing antimicrobial release profiles to be tailored for a given application and providing for sustained antimicrobial activity over time. More particularly, the invention relates to polymer compositions containing colloids comprised of salts of one or more oligodynamic metal, such as silver. The process of the invention includes mixing a solution of one or more oligodynamic metal salts with a polymer solution or dispersion and precipitating a colloid of the salts by addition of other salts to the solution which react with some or all of the first metal salts.

# POLYMER COMPOSITIONS CONTAINING COLLOIDS OF SILVER SALTS

## FIELD OF THE INVENTION

The present invention relates generally to polymer compositions and their use for making or coating articles, such as medical devices. More specifically the invention relates to antimicrobial compositions containing a polymer and oligodynamic salts.

## BACKGROUND OF THE INVENTION

For many years silver and silver salts have been used as antimicrobial agents. An early medicinal use of silver was the application of aqueous silver nitrate solutions to prevent eye infection in newborn babies. Silver salts, colloids, and complexes have also been used to prevent and to control infection. For example, colloidal metallic silver has been used topically for conjunctivitis, urethritis, and vaginitis.

Other metals, such as gold, zinc, copper, and cesium, have also been found to possess antimicrobial properties, both alone and in combination with silver. These and other metals have been shown to provide antimicrobial behavior even in minute quantities, a property referred to as "oligodynamic."

Additionally, silver is known for antimicrobial use with medical devices, such as catheters, cannulae, and stents. One conventional approach for obtaining antimicrobial medical devices is the deposition of metallic silver directly onto the surface of the substrate, for example, by vapor coating, sputter coating, or ion beam coating. However, these noncontact deposition coating techniques suffer many drawbacks. These drawbacks include poor adhesion, lack of coating uniformity, and the need for special processing conditions, such as preparation in darkness due to the light sensitivity of some silver salts. One particular drawback of these coatings is that the processes by which the coatings are formed do not adequately coat hidden or enclosed areas, such as the interior lumen of a catheter or stent. Additionally, these methods produce coatings that are very much like metallic silver in that they do not release silver from the coating and require contact with the coating to provide antimicrobial action. Though high concentrations of silver may be deposited on the substrate, very little free ionic silver is released on exposure to aqueous fluid. As a result, these coatings provide only limited antimicrobial activity. They essentially retard colonization of microbial agents on the surface of the device. However, because they do not release sufficient silver ions into aqueous fluids, they offer little or no protection from bacteria carried into the body upon insertion of the device and do not inhibit infection in the surrounding tissue.

Another method of coating silver onto a substrate involves deposition or electrodeposition of silver from solution. Drawbacks of these methods include poor adhesion, low silver pick-up on the substrate, the need for surface preparation, and 5 high labor costs associated with multistep dipping operations usually required to produce the coatings. Adhesion problems have been addressed by inclusion of deposition agents and stabilizing agents, such as gold and platinum metals, or by forming chemical complexes between a silver compound and the substrate 10 surface. However, inclusion of additional components increases the complexity and cost of producing such coatings.

With many medical devices, it is preferred to have a lubricious coating on the device. Lubricious coatings aid device insertion, reduce the trauma to tissue, and reduce the adhesion of 15 bacteria. Another drawback to conventional methods which apply silver and other metals directly onto the surface of a medical device for which a lubricious coating is also desired is that a second, lubricious coating must be applied to the device over the antimicrobial coating, adding to manufacturing cost and time.

20 Some of these coatings release, to varying degrees, silver ions into the solution or tissue surrounding the substrate. However, activation of such coatings often requires conditions that are not suitable for use with medical implants, such as catheters, stents, and cannulae. These conditions include abrasion 25 of the coating surface, heating to a temperature above 180°C,

contact with hydrogen peroxide, and treatment with an electric current.

Another conventional approach for obtaining antimicrobial medical devices is the incorporation of silver, silver 5 salts, and other antimicrobial compounds into the polymeric substrate material from which the article is formed. An oligodynamic metal may be physically incorporated into the polymeric substrate in a variety of ways. For example, a liquid solution of a silver salt may be dipped, sprayed or brushed onto 10 the solid polymer, for example, in pellet form, prior to formation of the polymeric article. Alternatively, a solid form of the silver salt can be mixed with a finely divided or liquefied polymeric resin, which is then molded into the article. Further, the oligodynamic compound can be mixed with monomers of the 15 material prior to polymerization.

There are several disadvantages to this approach. One such disadvantage is that larger quantities of the oligodynamic material are required to provide effective antimicrobial activity at the surface of the device. A second 20 disadvantage is that it is difficult to produce articles that allow for the release of the oligodynamic material because most device polymers absorb little, if any, water to aid in the diffusion and release of the oligodynamic material, resulting in articles that provide only a limited antimicrobial effect.

Yet another approach for obtaining antimicrobial medical devices is the incorporation of oligodynamic agents into a polymeric coating which is then applied to the surface of the article. Typically, an oligodynamic agent is incorporated into the 5 coating solution in the form of a solution or a suspension of particles of the oligodynamic agent. Problems associated with this approach include poor adhesion of the coating to the substrate, settling and agglomeration of the oligodynamic particles, and inadequate antimicrobial activity over time.

10 Settling of particles of the oligodynamic agent occurs as a result of the size and density of the particles. Settling of the particles from such solutions can cause unpredictable changes in the concentration of the oligodynamic agent in the composition. These changes in ion concentration result in several drawbacks to 15 producing commercial products. First, unpredictable changes in the concentration of the oligodynamic agent make it difficult to produce a composition having a specific concentration of antimicrobial ions and, thus, a particular effectiveness. Additionally, these changes make it difficult to produce multiple 20 batches of the composition having the same antibacterial concentration. Further, the concentration of the antimicrobial ions can affect other properties of the composition, such as its adhesive and lubricious properties. Consistency of antimicrobial activity is essential in the production of medical devices.

Another problem associated with particle suspensions is agglomeration of the particles. Particle agglomeration produces larger particle sizes which increases settling of particles from solution. Additionally, the agglomeration of particles in 5 suspensions and coating solutions can produce particles in the coating that are large enough to be noticeable to the touch on the coated surface. Articles produced using such coatings have decreased patient comfort and, therefore, are undesirable.

Many researchers have attempted to overcome these 10 problems. For example, U.S. Patent No. 4,592,920 to Murtfeldt *et al.* discloses a process that attempts to overcome the settling and agglomeration problems in the art through the use of a comminuted metal having a particle size of 30 microns or less. The coating of the *Murtfeldt* patent, however, exhibits several 15 disadvantages. For example, the *Murtfeldt* coating exhibits poor adhesion which is overcome by the use of the following methods. First, the *Murtfeldt* patent recommends pretreatment of the catheter to leach undesirable compounds that interfere with the bonding of the coating to the surface of the catheter. Second, the 20 *Murtfeldt* patent recommends the use of a bridging compound, or primer, to attach the coating to the surface of the catheter to increase adhesion. This adds an additional manufacturing step to the fabrication of a coated device. In addition to these disadvantages, it is likely that the process used to manufacture 25 and coat the catheters in *Murtfeldt* will result in settling and

agglomeration problems even with the use of silver having smaller particle sizes.

U.S. Patent No. 4,849,223 to Pratt *et al.* attempts to overcome settling and agglomeration of the particles in his invention by using solutions that contain high concentrations of polymer or monomer solids and are, thus, viscous. Suspending particles in high viscosity coating solutions containing high polymer solids is a common method for reducing settling and agglomeration of the particles. The coatings made by this method are usually very thick and, as a result, are often not uniform. Thick coatings are also more costly, dry more slowly than thin coatings, and are more difficult to manufacture. The coatings of the Pratt patent also exhibit poor adhesion. To increase adhesion, the Pratt patent recommends using coating materials which are similar to the substrate to be coated, pretreating the surface of the substrate before the coating composition is applied, or applying an additional coating layer between the substrate and the coating.

U.S. Patent No. 5,019,096 to Fox, Jr. *et al.* discloses a method for increasing the antibacterial activity of silver by incorporating a synergistic amount of chlorhexidine and a silver salt in a matrix-forming polymer. The polymer is such that it allows for release of the antimicrobial agent over an extended period of time. Fox, however, relies on dispersion of silver

particles into coating solutions and will be susceptible to problems associated with particle settling and agglomeration.

U.S. Patent No. 4,677,143 to Laurin *et al.* discloses a method to enhance release of the antimicrobial metal ions from the 5 surface of a device by incorporating the antimicrobial metal into a binder having a low dielectric constant that coats or forms the device. The nature of the binder allows the particles to form chain-like structures among themselves. These chain-like structures allow the surface particles to dissolve to provide an 10 initial dose of the antimicrobial agent and to create a pathway for interior particles to come to the surface to provide additional doses of the antimicrobial agent over time. *Laurin*, however, also relies on dispersion of silver particles into coating solutions and 15 is susceptible to problems associated with particle settling and agglomeration.

U.S. Patent No. 4,933,178 to Capelli discloses a polymer coating containing an oligodynamic metal salt of a sulfonylurea. The *Capelli* patent attempts to improve the solubility and stability of the antimicrobial metal in the coating and to 20 provide for the sustained release of the antimicrobial agent by adding a carboxylic acid to the coating composition. The particular carboxylic acids and the proportions in which they are mixed determine the rate of release of the antimicrobial agent from the polymer coating composition.

U.S. Patent No. 5,848,995 to Walder discloses the solid phase production of polymers containing AgCl as an antimicrobial agent. In the *Walder* process, solid polymer pellets are first soaked in a solution of silver nitrate which is absorbed into the 5 pellets. The pellets are then rinsed, dried, and soaked in a solution of a sodium chloride. The chloride ions of the salt are absorbed into the polymer matrix of the pellets where they react with the silver nitrate to form silver chloride. The pellets are then rinsed, dried, and melt processed. The compositions of the *Walder* 10 patent are limited to hydrophilic polymers, must be thermoformed, and do not contain other silver salts to provide multiple release rates, or other oligodynamic or medicinal agents to enhance antimicrobial effectiveness.

Therefore, there is a need in the art to provide a 15 method for rendering articles, such as medical devices, resistant to infection, not only on the surface of the article but also in the surrounding tissue. There is also a need in the art for compositions which can be incorporated into articles to provide antimicrobial activity. Further, there is a need for compositions 20 which can be employed as coatings for articles that exhibit improved adhesion. There is also a need for compositions that overcome the solubility, settling, and agglomeration problems of conventional oligodynamic compositions, and exhibit enhanced, sustained release of oligodynamic agents.

## SUMMARY OF THE INVENTION

Stated generally, the present invention comprises antimicrobial compositions which in a first aspect provide the advantage of reduced settling and agglomeration by producing a 5 minimal particle size of the oligodynamic salts in the compositions. The use of colloids in the compositions also permits incorporation of higher quantities of antimicrobial ions without the difficulties associated with the suspensions used in the prior art.

In another aspect, the compositions of the present 10 invention provide the advantage of varying release kinetics for the active oligodynamic ions due to the different water solubilities of the different salts in the compositions. These varying release kinetics allow for an initial release of oligodynamic ions that provides antimicrobial activity immediately upon insertion, 15 followed by a continual, extended release of the oligodynamic ions from the composition, resulting in sustained antimicrobial activity over time.

Stated somewhat more specifically, the present invention relates in one aspect to compositions that comprise a 20 polymer and a colloid containing salts of one or more oligodynamic agents. In one disclosed embodiment, the polymer is a hydrophilic polymer. In another disclosed embodiment, the polymer is a hydrophobic polymer, while in yet another

embodiment, the polymer is a combination of these two types of polymers.

In one disclosed embodiment, the invention comprises one or more salts of silver as the oligodynamic agent. In another 5 embodiment, the composition optionally contains additional salts of other oligodynamic metals, such as zinc, gold, copper, cerium and the like. In yet another embodiment, the composition optionally comprises additional salts of one or more noble metals to promote galvanic action. In still another embodiment, the 10 composition optionally comprises additional salts of platinum group metals such as platinum, palladium, rhodium, iridium, ruthenium, osmium, and the like. The compositions optionally contain any other components that provide beneficial properties to the composition or improve the antimicrobial effectiveness of 15 the composition.

In a second aspect, the present invention relates to a process for producing these antimicrobial compositions. The process comprises the formation of colloids of oligodynamic agents in solutions, dispersions, or combinations of polymers 20 solutions and dispersions. The terms "polymer composition" and "polymer solution" are used interchangeably throughout the specification and claims and both means any polymer solution, dispersion, or combination of polymer solutions and dispersions. The colloid can be formed first and then added to the polymer 25 composition or can be formed *in situ* in the polymer composition.

Preferably, the colloid is formed *in situ* in the polymer composition.

The process of forming the colloids comprises, for example, combining two or more salts, wherein at least one of the 5 salts is the salt of an oligodynamic agent. These salts will be referred to herein as salt A and salt B. Salt A comprises one or more oligodynamic agents. Salt B comprises one or more salts that can react with salt A to form a colloid. Salts A and B can be combined in any amount and in any order. However, it is 10 preferred that salt A be present in a stoichiometric amount or in excess when compared to salt B.

Optionally, additional components can be added to the antimicrobial compositions of the present invention. These components include, but are not limited to, additional 15 oligodynamic agents, additional soluble salts, salts which provide galvanic action, and any other components which provide the compositions with beneficial properties or enhance the antimicrobial activity of the compositions. Such components include, but are not limited to, antimicrobial agents, antibiotics, 20 and other medicinal agents.

In one disclosed embodiment, the antimicrobial composition of the invention is produced by forming a solution, dispersion, or combination of solutions and dispersions of one or more polymers. Next, a solution comprising salt A is added to the

polymer composition. Then, a solution comprising salt B is added to the polymer composition to precipitate fine colloidal salt(s) of the oligodynamic agent(s). Where the oligodynamic agent is a metal salt, the metal cation of salt A reacts with the anion of salt B 5 to form a less soluble salt which precipitates as a fine colloid. Salt B is added to the polymer composition in an amount sufficient to react with some or all of salt A. Optionally, other salts are then added in amounts to react with some or all of the remaining amount of salt A.

10 In another disclosed embodiment, salt B is added to the polymer composition, followed by the addition of an excess or stoichiometric amount of salt A. In yet another embodiment, salts A and B can be combined to form a colloid which is then added to the polymer composition.

15 The final polymer composition formed by these processes contains one or more colloidal salts, composed of the oligodynamic cations of salt A and the anions of salt B, and one or more soluble salts, composed of the anions of salt A and the cations of salt B.

20 In another aspect, the present invention relates to an article of manufacture which comprises a substrate and the antimicrobial compositions of the present invention. In a disclosed embodiment, the composition is employed in the manufacture of the article itself. Thus, the final article is composed of one or more

of the compositions of the present invention, alone or in admixture with other polymeric components. In another disclosed embodiment, the composition is applied to a preformed article as a coating. The coated article may be produced, for example, by 5 dipping the article into the composition or by spraying the article with the composition and then drying the coated article. In a preferred embodiment, the compositions are used to coat medical devices.

## DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

### 10 THE COMPOSITION

In a first aspect, the present invention provides antimicrobial compositions. The compositions comprise a polymer and a colloid comprised of the salts of one or more oligodynamic agents. The term "oligodynamic agents" as used in the present 15 invention refers to any compound that can provide antimicrobial activity, even when present in small quantities.

Any polymer may be employed in the present invention, including hydrophilic polymers, hydrophobic polymers, and mixtures of these two types of polymers. The use of 20 hydrophilic polymers is preferred because such polymers have additional benefits. These benefits include increased lubricity for patient comfort, increased absorption of aqueous fluids from the body which aids in the release of oligodynamic ions from the

composition, inhibition of bacterial attachment, and improved solubility for some metal salts. Hydrophilic polymers best suited to the invention are those that are soluble in water or in organic solvents containing water. The ability to add water to the 5 polymer composition without precipitating the polymer allows the addition of water-soluble salts directly to the coating composition. Water facilitates the formation of salt colloids within the polymer composition. For this reason, it is preferred that the polymer solution contain from 1 to 50% water by weight, more preferably 10 from 5 to 30% water.

However, the use of water is not limiting, as salt colloids can also be formed using alcohols, organic solvents, or both that contain little or no water. The use of alcohols and organic solvents, containing from 0 to 1% water are preferred 15 when hydrophobic polymers are employed in the present invention.

Examples of hydrophilic polymers which may be used to form the compositions include, but are not limited to, polyurethanes, including polyether polyurethanes, polyester 20 polyurethanes, polyurethaneureas, and their copolymers; polyvinylpyrrolidones; polyvinyl alcohols; polyethylene glycols and their copolymers; polypropylene glycols and their copolymers; polyoxyethylenes and their copolymers; polyacrylic acid; polyacrylamide; carboxymethyl cellulose; cellulose and its 25 derivatives; dextrans and other polysaccharides; starches; guar;

xantham and other gums and thickeners; collagen; gelatins; and other biological polymers. Preferred hydrophilic polymers are polyurethanes and polyurethane copolymers, such as polyether polyurethaneurea.

5 Examples of hydrophobic polymers best suited for use in the present invention include, but are not limited to, polytetrafluoroethylene, polyvinyl chloride (PVC), polyvinylacetate, poly(ethylene terephthalate), silicone, polyesters, polyamides, polyureas, styrene-block copolymers, polymethyl 10 methacrylate, acrylic-butadiene-styrene copolymers, polyethylene, polystyrene, polypropylene, natural and synthetic rubbers, acrylonitrile rubber, and mixtures and copolymers of any of the above. The preferred hydrophobic polymer depends upon the substrate to be coated. Hydrophobic polymers that are chemically 15 similar or identical to the substrate are advantageously used alone or in combination with hydrophilic polymers to form coatings that enhance adhesion of the coating to the substrate.

The colloid of the present invention comprises one or more oligodynamic salts. In the discussion of the process below, 20 these salts are referred to as salt A. In a preferred embodiment, the oligodynamic salts comprise one or more salts of oligodynamic metals. The salts may be different salts of the same oligodynamic metal or may be salts of different oligodynamic metals. Oligodynamic metals useful in the present invention include, but 25 are not limited to, silver, platinum, gold, zinc, copper, cerium,

gallium, osmium, and the like. The preferred oligodynamic metal is silver.

Salts of other metals may be employed to form the colloid. In the discussion of the process below, these salts are referred to as salt B. These salts contain cationic ions that include, but are not limited to, calcium, sodium, lithium, aluminum, magnesium, potassium, manganese, and the like, and may also include oligodynamic metal cations such as copper, zinc, and the like. These salts contain anions that include, but are not limited to, acetates, ascorbates, benzoates, bitartrates, bromides, carbonates, chlorides, citrates, folates, gluconates, iodates, iodides, lactates, laurates, oxalates, oxides, palmitates, perborates, phenosulfonates, phosphates, propionates, salicylates, stearates, succinates, sulfadiazines, sulfates, sulfides, sulfonates, tartrates, thiocyanates, thioglycolates, thiosulfates, and the like.

The compositions of the present invention can also contain additional antimicrobial materials. For example, the compositions can contain salts of metals that enhance the antimicrobial effect of the oligodynamic metal, such as the platinum group metals, or other metals that promote galvanic action.

The compositions of the present invention can also contain any combination of additional medicinal compounds. Such medicinal compounds include, but are not limited to, antimicrobials, antibiotics, antifungal agents, antiviral agents,

antithrombogenic agents, anesthetics, anti-inflammatory agents, analgesics, anticancer agents, vasodilation substances, wound healing agents, angiogenic agents, angiostatic agents, immune boosting agents, growth factors, and other biological agents.

5 Suitable antimicrobial agents include, but are not limited to, biguanide compounds, such as chlorhexidine and its salts; triclosan; penicillins; tetracyclines; aminoglycosides, such as gentamicin and Tobramycin™; polymyxins; rifampicins; bacitracins; erythromycins; vancomycins; neomycins;

10 chloramphenicols; miconazole; quinolones, such as oxolinic acid, norfloxacin, nalidixic acid, pefloxacin, enoxacin, and ciprofloxacin; sulfonamides; nonoxynol 9; fusidic acid; cephalosporins; and combinations of such compounds and similar compounds. The additional antimicrobial compounds provide for enhanced

15 antimicrobial activity.

The compositions can also contain auxiliary components. Examples of such auxiliary components include, but are not limited to, viscosity and flow control agents, antioxidants, conventional pigments, air release agents or defoamers, and

20 discolorants. The composition may also contain conventional dyes and pigments to impart color or radiopacity or to enhance the aesthetic appearance of the compositions. The compositions can also contain additional lubricating agents and other additives that enhance patient comfort and tissue health.

While not wishing to be bound by the following mechanism, it is believed that many of the advantageous properties of the present compositions result from the differences in the solubility of the different metal salts present in the colloid.

- 5 These differing solubilities of the metal salts in the colloid provide varying release kinetics for the active oligodynamic metal(s). For example, with a medical device composed of, or coated with, the compositions of the present invention, those salts that have high water solubility will be released from the coating rather quickly,
- 10 providing a high initial dose of antimicrobial activity to kill bacteria introduced upon insertion of the device in the patient. This initial dose is sometimes referred to as "quick kill," and this antimicrobial activity is identified by the ability of a coated device or composition to create zones of no bacterial growth around the
- 15 device or composition when it is placed in a bacterial culture. This test is known as a "zone of inhibition" assay. Those salts having lower water solubilities will be released more slowly from the composition, resulting in a sustained or extended antimicrobial activity over time.

- 20 Selection of salts having varying degrees of solubility in the composition allows tailoring of the composition to the specific application of the article comprising the composition. Specifically, compositions of the invention can be tailored to kill bacteria introduced during the insertion of a medical device, both
- 25 on the surface of the device and in the surrounding fluid and

tissue, by both the quick release of antimicrobial metal salts, followed by prolonged inhibition of bacterial migration and growth by the slower release of less soluble antimicrobial metal salts over an extended period of time. The ability to tailor the 5 release of the oligodynamic agent is advantageous over conventional antimicrobial compositions, as it provides for both immediate and sustained antimicrobial activity.

Another advantage of the compositions of the present invention is that the formation of colloids within the polymer 10 composition produces ultra-fine particles that possess a minimal particle size for the metal salts. This minimal particle size retards settling and agglomeration. The use of colloids in the composition also permits incorporation of higher quantities of antimicrobial metal without the difficulties associated with the suspensions used 15 in the prior art.

By reducing or eliminating the problems associated with conventional antimicrobial polymer compositions, the present invention provides reproducible compositions having a specific antimicrobial ion concentration with a specific antimicrobial ion 20 release profile that can be tailored through the specific salt combinations selected to provide optimum antibiotic activity over an extended period of time. For example, compositions of the invention can be tailored to release the bulk of their oligodynamic agents within 5 days for a medical device with a short term use in

the body, such as a wound drain, or within 30 days for a device with a longer term use, such as a foley catheter.

The tailored delivery embodiment of the invention will now be further described in terms of a polyurethane composition 5 containing a colloid of specific silver salts. It is to be understood that this is simply an example of one embodiment of the invention and that one of skill in the art, based upon the present disclosure, could pick and choose salts having differing solubilities to provide a composition having a suitable release profile for a particular 10 purpose.

A coating solution is formed from a 4.7% solution of a polyether polyurethane-urea block copolymer in a mixture of THF/alcohol in a 75/25 ratio by weight. A sufficient quantity of 10% silver nitrate ( $\text{AgNO}_3$ ) solution in water is added to the TPH 15 copolymer solution to produce a final silver concentration of approximately 15%, based on the weight of coating solids in the solution.

Aqueous solutions of sodium chloride, zinc iodide, sodium citrate, sodium acetate, and sodium lactate (each 1.0% 20 solutions) are added to the copolymer solution in sufficient amounts for each salt to react with 15% of the silver nitrate present in the composition. Colloids of silver chloride, silver iodide, silver citrate, silver acetate, and silver lactate are formed in the final coating composition. The coating composition also

contains 25% unreacted soluble silver nitrate, as well as the silver nitrate and zinc nitrate salt products. The differences in the solubility of the different salts in the composition will result in different and prolonged rates of release of the oligodynamic silver 5 in the coating composition when a device coated with the composition is exposed to body fluid.

Silver nitrate is the most soluble of the salts present in the composition and will be released rapidly upon initial exposure of the coating to body fluid. Sodium lactate, which has a lower 10 solubility than silver nitrate but a higher solubility than the other salts present, will be released next. Then, the silver acetate, followed by the silver citrate, and then the silver chloride, and, lastly, the silver iodide will be released from the coating composition based upon their relative solubilities.

15 The initial release and the duration of release of the oligodynamic agents from the composition depends upon several factors. These factors include the relative water solubilities of the particular salts formed in the colloid and the concentration of the salts in the colloid. This release can range, for example, from a 20 few days to several months, and can be tailored through the choice and number of salts formed in the composition for the intended purpose of the device to be coated.

The compositions of the invention can also be tailored to provide other desired properties, such as surface lubricity.

Further, the compositions may contain other medicinal or otherwise beneficial agents.

#### PROCESS FOR PREPARING THE COMPOSITION

In a second aspect, the present invention relates to a 5 process for producing the compositions of the invention. In general terms, the process comprises the formation of colloids of oligodynamic agents in polymer solutions. The colloid can be formed first and then added to the polymer composition or can be formed *in situ* in the polymer composition. Preferably, the 10 colloid is formed *in situ* in the polymer composition.

The process of forming the colloids comprises, for example, combining two or more salts, wherein at least one of the salts is the salt of an oligodynamic agent. These salts will be referred to as salt A and salt B. Salt A comprises one or more 15 oligodynamic agents. Salt B comprises one or more salts that can react with salt A to form a colloid. Salts A and B can be combined in any amount and in any order. However, it is preferred that salt A be present in a stoichiometric amount or in excess when compared to salt B.

20 Optionally, additional components can be added to the compositions. These components include, but are not limited to, additional oligodynamic agents, additional soluble salts, salts which provide galvanic action, and any other components which

provide the compositions with beneficial properties or enhance the antimicrobial activity of the compositions. Such components include, but are not limited to, antimicrobial agents, antibiotics, and other medicinal agents.

5 In one disclosed embodiment, the composition is produced by forming a solution, dispersion, or combination of solutions and suspensions of one or more polymers. Next, a solution comprising salt A is added to the polymer composition. Then, a solution comprising salt B is added to the polymer  
10 10 composition to precipitate fine colloidal salt(s) of the oligodynamic agent(s) of salt A. Where the oligodynamic agent is a metal salt, the metal cation of salt A reacts with the anion of salt B. Salt B is added to the polymer composition in an amount sufficient to react with some or all of salt A. Optionally, other salts are then added  
15 15 in amounts to react with some or all of the remaining amount of salt A.

In another disclosed embodiment, salt B is added to the polymer composition, followed by the addition of an excess or stoichiometric amount of salt A. In yet another embodiment, salts  
20 20 A and B can be combined to form a colloid which is then added to the polymer composition.

The final polymer composition formed by these processes contains one or more colloidal salts, composed of the oligodynamic cations of salt A and the anions of salt B, and one or

more soluble salts, composed of the anions of salt A and the cations of salt B. Additionally, other salts may be added to the composition that do not react in solution but provide some beneficial effect such as stabilization of the colloid, modification of

5 antimicrobial ion release rate, promotion of galvanic action, increase in antimicrobial effectiveness, or enhancement of biocompatibility. Further, other compounds may be added to the composition, including, but not limited to, medicinal agents, lubricants, nutritional agents, antioxidants, dyes and pigments,

10 and other additives.

As noted above, any polymer can be used to form the compositions of the present invention. When hydrophilic polymers are used, it is preferable that the polymers be soluble in water or in organic solvents containing some water. The ability to

15 add water to the polymer composition without precipitating the polymer allows the addition of water-soluble salts directly to the coating composition. The use of water in the polymer composition increases the solubility of the salts, resulting in the formation of finer, more stable colloids. However, it takes longer for the

20 coating compositions to dry when the water content is very high. For this reason, the preferred amount of water in the hydrophilic polymer compositions is about 50% or less. Such concentrations provide for faster drying times while maintaining the beneficial properties provided by the water in the composition.

In contrast, when hydrophobic polymers are used either alone or in combination with hydrophilic polymers, it is desirable to limit the amount of water present in the composition to avoid precipitation of the hydrophobic polymer with the 5 colloid. In such instances the amount of water present in the polymer composition is preferably 1% or less. While it is possible to practice the invention in the absence of water in the composition, it is preferable to have some water present. Thus, when hydrophobic polymers are employed in the present 10 invention, the preferred water content of the polymer compositions is between about 0.1% and 1% by weight. It is advantageous to employ salts that are soluble in alcohols or organic solvents when hydrophobic polymers employed.

Examples of water-soluble silver salts suitable for use 15 in the present invention, include, but are not limited to, silver nitrate, silver acetate and silver lactate. Examples of silver salts which are soluble in alcohols and organic solvents include, but are not limited to, silver nitrate, sodium iodide, sodium lactate, sodium propionate, sodium salicylate, zinc chloride, zinc acetate, zinc 20 salicylate, gold trichloride, gold tribromide, palladium chloride and hydrogen-hexachloroplatinate.

Examples of alcohols that are useful in the present 25 invention include, but are not limited to, methanol, ethanol, propanol, isopropanol, and butanol. Examples of organic solvents that can be used to form solutions of the oligodynamic salts

include, but are not limited to, acetone, tetrahydrofuran (THF), dimethylformamide (DMF), dimethylsulfoxide (DMSO), and acetonitrile. These organic solvents are especially useful when they contain a small amount of water.

5 It is also possible to prepare polymer compositions from supercritical fluids. The most common of these fluids is liquefied carbon dioxide.

In a preferred embodiment, the polymer composition in which the colloid is formed is a hydrophilic polyether 10 polyurethaneurea. This polymer is a substantially noncovalently crosslinked reaction product of one or more diols, water and an organic diisocyanate. The urea segments of the polymer provide improved strength, increased viscoelasticity, and decreased water absorption. These polymers typically absorb water in amounts 15 from 50 to 100% their weight while remaining strong and elastic.

Diols useful in the formation of these polymers include, but are not limited to, medium and long chain poly(oxyethylene) glycols having a number average molecular weights between 250 and 20,000. Examples of such diols are "Carbowax" compounds 20 sold by Union Carbide.

Organic diisocyanates useful to form these polymers include, but are not limited to, tetramethylene diisocyanate, hexamethylene diisocyanate, trimethylhexamethylene diisocyanate, dimer acid diisocyanate, isophorone diisocyanate,

diethylbenzene diisocyanate, decamethylene 1,10-diisocyanate, cyclohexylene 1,2-diisocyanate, cyclohexylene 1,4-diisocyanate, 5 methylene bis(cyclohexyl-4-isocyanate), 2,4- and 2,6-tolylene diisocyanate, 4,4-diphenylmethane diisocyanate, 1,5-naphthaliene diisocyanate, dianisidine diisocyanate, tolidine diisocyanate, 10 xylylene diisocyanate, and tetrahydronaphthalene-1,5-diisocyanate.

In another preferred embodiment, the polymer coating composition comprises a combination of a hydrophilic 10 polyurethane, a polymer that is similar or identical to the polymer substrate to be coated, and, optionally, other polymers which aid coating adhesion and physical properties. Antimicrobial salt colloids are prepared in this composition as disclosed previously, with the exception that, depending on the second polymer used, 15 some or all of the water used to prepare salt solutions can be replaced with alcohols or other organic solvents to prevent precipitation of the second polymer. Another exception is that the salts elected must be soluble in solvents compatible with those in which the polymers are soluble. As an example of this preferred 20 embodiment, a solution of a hydrophilic polyether polyurethaneurea in THF can be combined with a solution of polyvinyl chloride (PVC) in methylene chloride in equal amounts. Then, silver nitrate can be dissolved in ethanol and added to the 25 solution without precipitation. Ethanol is used to dissolve the silver nitrate instead of water because PVC has a tendency to

precipitate when water is added to the solution. Finally, a dilute solution of zinc chloride in ethanol/water can be slowly added to the polymer composition to produce a fine silver chloride colloid without precipitation of the PVC. The final concentration of  
5 water in the coating is less than 1%. The coating solution is then used to dip-coat PVC catheters. The finished coating is well adhered, durable, lubricious when wetted, and contains colloidal antimicrobial salts.

In another embodiment, the polymer composition  
10 comprises a hydrophilic polymer as defined in commonly owned copending application Serial No. 09/189,240, filed November 10, 1998, herein incorporated by reference. In general, the polymer is a polyurethane-urea-silane copolymer prepared from the following ingredients: (1) one or more polyisocyanate, (2) one or  
15 more lubricious polymer having at least two functional groups, which may be the same or different and are reactive with an isocyanate functional group, and (3) one or more organo-functional silanes having at least two functional groups, which may be the same or different and are reactive with an isocyanate functional group and with a silicone rubber substrate. While these copolymers may be prepared in a variety of ways, preferably they may be prepared by first forming a prepolymer from the polyisocyanate(s) and lubricious polymer(s) followed by reaction with the organo-functional silane(s). A catalyst is optionally  
20 employed during reaction of the isocyanate with the polyol.  
25

Isocyanates useful to form these polymers include, but are not limited to, 4,4'-diphenylmethane diisocyanate and position isomers thereof, 2,4- and 2,6- toluene diisocyanate (TDI) and position isomers thereof, 3,4-dichlorophenyl diisocyanate, 5 dicyclohexylmethane-4,4'-diisocyanate (HMDI), 4,4'-diphenylmethane diisocyanate (MDI), 1,6-hexamethylene diisocyanate (HDI) and position isomers thereof, isophorone diisocyanate (IPDI), and adducts of diisocyanates, such as the adduct of trimethylolpropane and diphenylmethane diisocyanate 10 or toluene diisocyanate.

Polyols useful to form these polymers include, but are not limited to, polyethylene glycols, polyester polyols, polyether polyols, castor oil polyols, and polyacrylate polyols, including Desmophen A450, Desmophen A365, and Desmophen A160 15 (available from Mobay Corporation), poly(ethylene adipates), poly(diethyleneglycol adipates), polycaprolactone diols, polycaprolactone-polyadipate copolymer diols, poly(ethylene-terephthalate)diols, polycarbonate diols, polytetramethylene ether glycol, ethylene oxide adducts of polyoxypolyethylene diols, and 20 ethylene oxide adducts of polyoxypolyethylene triols.

Catalysts useful to form these polymers include, but are not limited to, tertiary amines, such as N,N-dimethylaminoethanol, N,N-dimethyl-cyclohexamine-bis(2-dimethyl aminoethyl) ether, N-ethylmorpholine, N,N,N',N",N"-pentamethyl-diethylene-triamine, and 25 1-2(hydroxypropyl)

imidazole, and metallic catalysts, such as tin, stannous octoate, dibutyl tin dilaurate, dioctyl tin dilaurate, dibutyl tin mercaptide, ferric acetylacetone, lead octoate, and dibutyl tin diricinoleate.

Silanes useful to form these polymers include, but are  
5 not limited to, N-beta-(aminoethyl)-gamma-aminopropyl-  
trimethoxy silane and diamino-alkoxysilanes, such as N-(2-  
aminoethyl)-3-aminopropylmethyl-dimethoxy silane.

These polymers preferably have from 7 to 12% by  
weight silane based upon the weight of the entire polymer. The  
10 preferred ratio of isocyanate functional groups to alcohol or other  
isocyanate reactive functional groups is from 1.1:1 to 2:1.  
Viscosity of the polymer solution is a function of molecular weight  
of the polymer and the solids content of the solution and is  
controlled by addition of solvent to the solution. The preferred  
15 copolymer solution for dip coating has a kinematic viscosity in the  
range of about 1.5 cS to about 20 cS (centistokes), and a solids  
content in a range of about 0.4 to about 5.

In yet another embodiment, the polymer composition  
comprises a solution of a hydrophilic polymer as defined in U.S.  
20 Patent No. 5,290,585, which is hereby incorporated by reference.  
The polymer is a polyurethane-polyvinyl pyrrolidone prepared by  
mixing the appropriate amounts of isocyanate, polyol, and  
polyvinyl pyrrolidone (PVP) stock solution. Additional solvents  
can be added to adjust the viscosity and solids content. Solids

content may be in the range of 0.4 to 15% by weight, depending on the solvent used and other considerations. The stoichiometric ratio of total NCO groups in the isocyanate to total OH groups in the polyol may vary from 0.75 to 3.0. Preferably, the isocyanate 5 has at least two NCO groups per molecule and the polyol has at least two OH groups per molecule. The ratio of polyurethane formed *in situ* to PVP ranges from 0.05 to 3.0 by weight.

The PVP employed to form these polymers preferably has a mean molecular weight from about 50,000 to 2.5 million 10 Daltons. Specific preferred PVP polymers are Kollidon 90, Luviskol K90, Luviskol K80, and Luviskol K60, all available from BASF Corp. (Parsippany, NJ) and Plasdone 90, PVP K90, and PVP K120, all available from GAF Corporation.

Isocyanates suitable to form these polymers include, 15 but are not limited to, polymethylenepolyphenyl isocyanate, 4,4'-diphenylmethane diisocyanate and position isomers thereof, 2,4-tolylene diisocyanate and position isomers thereof, 3,4-dichlorophenyl diisocyanate, isophorone isocyanate, and adducts or prepolymers of isocyanates, such as the isocyanate prepolymer 20 available as Vorite 63 from CasChem, Inc. (Bayonne, NJ). Other examples of polyisocyanates useful in the present invention are those listed in ICI Polyurethanes Book, by George Woods, published by John Wiley and Sons, New York, New York (1987).

Polyols useful to form these polymers include, but are not limited to, polyester polyols, polyether polyols, modified polyether polyols, polyester ether polyols, castor oil polyols, and polyacrylate polyols, including Desmophen A450, Desmophen 5 A365, and Desmophen A160 available from Mobay Corporation (Pittsburgh, PA). Preferred polyols include castor oil and castor oil derivatives, such as DB oil, Polycin-12, Polycin 55, and Polycin 99F available from CasChem, Inc. Preferred diols include, but are not limited to, Desmophen 651A-65, Desmophen 1300-75, 10 Desmophen 800, Desmophen-550 DU, Desmophen-1600U, Desmophen-1920D, and Desmophen-1150, available from Mobay Corporation, and Niax E-59 and others available from Union Carbide (Danbury, Connecticut).

15 Suitable solvents for use in the formation of these polymers are those which are capable of dissolving the isocyanate, the polyol, and the polyvinyl pyrrolidone without reacting with any of these components. Preferred solvents include, but are not limited to, methylene chloride, dibromomethane, chloroform, dichloroethane, and dichloroethylene.

20 When a composition containing this polymeric solution is to be used as a coating, the coating is cured, after application to the substrate, at a temperature in the range of approximately 75°F to approximately 350°F for a period in the range of about 2 minutes to about 72 hours.

The process of the invention will now be further described in terms of the formation of a colloid of silver chloride from silver nitrate and sodium chloride in a polyurethane polymer coating solution. It is to be understood that this is simply an 5 example of one preferred embodiment of the invention and that any polymer or combination of polymers and any mixture of salts that will form a colloid within the polymer solution can be employed in the present invention.

First, a 4.7% solution of a polyether polyurethane-urea 10 block copolymer is prepared in a mixture of THF/ethanol in a 75/25 ratio by weight. A sufficient quantity of 10% silver nitrate ( $\text{AgNO}_3$ ) solution in water is added to the TPH copolymer solution to produce a final silver concentration of approximately 15%, based on coating solids in the solution. An aqueous solution 15 of 1.0% sodium chloride ( $\text{NaCl}$ ) is then slowly added to the solution with stirring in an amount sufficient to react with 50% of the  $\text{AgNO}_3$ . The  $\text{NaCl}$  reacts with the  $\text{AgNO}_3$  to produce a colloidal suspension of the poorly water soluble salt,  $\text{AgCl}$ , and the soluble salt,  $\text{NaNO}_3$ , from half of the  $\text{AgNO}_3$ . The amount of 20 water in the final coating solution is about 30% of the total solvent weight. The final polymer concentration in the coating solution is 3.3%, based upon solvent and polymer weights.

A 16 Fr latex Foley catheter can then be coated with the composition by dipping it into the composition solution, 25 withdrawing it at a controlled rate and drying it using standard

methods. The finished coating contains both the water soluble, and therefore fast releasing, AgNO<sub>3</sub>, and the water insoluble, and therefore slow releasing, AgCl.

#### USES OF THE COMPOSITION

5           In a third aspect, the present invention relates to an article of manufacture. In a preferred embodiment, the antimicrobial composition can be used as a coating on a preformed article to provide antimicrobial activity to the surface of the article and to the environment surrounding the article through the

10 continual release of oligodynamic ions. Any article can be coated with the antimicrobial compositions of the present invention. The composition is particularly suited for the production of medical devices, which include, but are not limited to, catheters, cannulae, stents, guide wires, implant devices, contact lenses, IUDs,

15 peristaltic pump chambers, endotracheal tubes, gastroenteric feeding tubes, arteriovenous shunts, condoms, oxygenator and kidney membranes, gloves, pacemaker leads, and wound dressings.

In a second embodiment, the antimicrobial composition

20 can be prepared as a high solids solution and used alone or mixed with other polymers to form the article itself. Polymers which are useful to form the articles of the invention include, but are not limited to, natural and synthetic rubber, especially latex rubber, acrylonitrile rubber, PVC plastisol, PVC, polyurethanes, silicone,

polycarbonates, acrylates, polyamides, polypropylenes, polyethylenes, polytetrafluoroethylenes, polyvinylacetate, poly(ethylene terephthalate), polyesters, polyamides, polyureas, styrene-block copolymers, polymethyl methacrylate, acrylic-  
5 butadiene-styrene copolymers, polystyrene, cellulose, and derivatives and copolymers of any of the above.

As nonlimiting examples, compositions of the invention can be admixed into latex rubber for fabrication of catheters, gloves, and other dipped latex products by standard form  
10 dipping methods, and vinyl plastisols can be mixed with compositions of the invention to provide dippable and castable antimicrobial PVC devices. Thus, the final article can be composed of one or more of the compositions of the present invention in admixture with other polymeric components.

15 Alternatively, compositions of the invention can be formulated into high solids coating compositions that can be used to dip-fabricate a variety of medical devices, such as catheters, stents, gloves, condoms, and the like.

By another method, compositions of the invention can  
20 be dried and melt processed, for example, by injection molding and extrusion. Compositions used for this method can be used alone or compounded with any other melt-processable material for molding and extrusion of antimicrobial articles.

The antimicrobial compositions of the invention are preferably coated onto preformed articles. When used as a coating, the compositions can be applied by any means, including those methods known in the art. For example, the compositions 5 can be brushed or sprayed onto the article, or the article can be dipped into the composition. For example, the article can be dipped into the antimicrobial polymer solution at a rate of about 10-80 inches per minute (ipm), preferably about 40 ipm. The article is allowed to remain in the antimicrobial polymer solution 10 for a period of about 0-30 seconds, preferably about 5-15 seconds. The article is then withdrawn at a rate of about 10-80 ipm, preferably about 15-30 ipm. Once the article has been coated with the copolymer of the invention, it is allowed to air dry for a period of at least about 10 minutes before drying is completed in 15 an oven for a period of about 5-60 minutes at a temperature in the range of about 40-100°C. Preferably, oven drying occurs for a period of about 15 minutes at a temperature of about 50°C. The coated article can optionally be dried with a hot air stream at a temperature in the range of approximately 40°C to approximately 20 100°C for a period of about 5-60 minutes to remove residual solvent.

This invention is further illustrated by the following examples, which are not to be construed in any way as imposing limitations upon the scope of the invention.

## EXAMPLES

Example 1

To form the coating solution, a 4.7% solution of a polyether polyurethane-urea block copolymer manufactured by 5 Tyndale Plains-Hunter Ltd. (TPH) was prepared in a mixture of THF/alcohol in a 75/25 ratio by weight. A sufficient quantity of 10% silver nitrate ( $\text{AgNO}_3$ ) solution in water was added to the TPH copolymer solution to produce a final silver concentration of approximately 15%, based on the weight of coating solids in the 10 solution. An aqueous solution of 1.0% sodium chloride ( $\text{NaCl}$ ) was added to the solution in an amount sufficient to react with 50% of the  $\text{AgNO}_3$  to produce a colloid of the poorly water soluble salt,  $\text{AgCl}$ , from half of the  $\text{AgNO}_3$  in the coating solution. The  $\text{NaCl}$  solution was added slowly to the polymer solution and 15 the solution began to turn cloudy with the formation of the fine colloidal  $\text{AgCl}$ . The amount of water in the final coating solution was about 30% of the total solvent weight. The final polymer concentration in the coating solution was 3.3%, based upon solvent and polymer weights.

20 A 16 Fr latex Foley catheter was then coated by dipping it into the coating solution, withdrawing it at a controlled rate to control the thickness of the coating and drying the catheter coating using standard methods. The finished coating contained both the water soluble, and therefore fast releasing,

AgNO<sub>3</sub> and the water insoluble, and therefore slow releasing, AgCl.

Example 2

The process of Example 1 was repeated, except that a  
5 1.0% solution of zinc chloride was used in place of the 1.0%  
solution of sodium chloride, resulting in the formation of a silver  
chloride colloid and zinc nitrate from half the silver nitrate in the  
coating solution. Zinc chloride was added in an amount of one  
half the amount of NaCl added in Example 1 because one mole of  
10 zinc chloride reacts with 2 moles of silver nitrate.

Example 3

The process of Example 1 was repeated, except that a  
1.0% solution of copper chloride was used in place of the 1.0%  
solution of sodium chloride, resulting in the formation of a silver  
15 chloride colloid and copper nitrate from half the silver nitrate in  
the coating solution. Copper chloride was added in an amount of  
one half the amount of NaCl added in Example 1 because one  
mole of copper chloride reacts with 2 moles of silver nitrate.

Example 4

20 The process of Example 1 was repeated, except that  
the 1.0% solution of sodium chloride was replaced with a 1.0%  
solution of sodium iodide, resulting in the formation of a silver

iodide colloid and sodium nitrate from half the silver nitrate in the coating solution. Silver iodide is a local antiinfective agent and has a lower water solubility than silver chloride, providing a slower releasing silver salt than silver chloride.

5 Example 5

The process of Example 1 was repeated, except that the 1.0% solution of sodium chloride was replaced with a 1.0% solution sodium propionate, resulting in the formation of a silver propionate colloid and soluble sodium nitrate, along with the 10 remaining silver nitrate in the solution. Silver propionate is a local antiinfective and is more water soluble than AgCl or AgI, providing a faster releasing salt than silver chloride or silver iodide.

Example 6

15 The process of Example 1 was repeated, except that the 1.0% solution of sodium chloride was replaced with a 1.0% solution of sodium lactate, resulting the formation of a silver lactate colloid and sodium nitrate, along with the remaining silver nitrate in the coating solution. Silver lactate is a local antiinfective 20 and is more water soluble than sodium propionate, AgCl or AgI, providing one of the fastest releasing silver salts, other than the soluble silver nitrate.

Example 7

The process of Example 1 was repeated, except that the 1.0% solution of sodium chloride was replaced with a solution of sodium acetate, resulting in the formation of a silver acetate colloid and sodium nitrate, along with the remaining silver nitrate in the solution. Silver acetate is a local antiinfective that is more water soluble than sodium propionate, silver chloride, or silver iodide, but less water soluble than silver lactate.

5

Example 8

10 The process of each of Examples 1, 2, 3, 4, 5, 6, and 7 was repeated, except that the salt solution was added in an amount sufficient to react with 75% of the  $\text{AgNO}_3$ .

Example 9

The process of each of Examples 1, 2, 3, 4, 5, 6, and 7  
15 was repeated, except that the salt solution was added in an amount sufficient to react with 100% of the  $\text{AgNO}_3$ .

Example 10

The process of each of Examples 1, 2, 3, 4, 5, 6, and 7 was repeated, except that the salt solution was added in an  
20 amount sufficient to react with 25% of the  $\text{AgNO}_3$ .

Example 11

The process of Example 1 was repeated, except that the NaCl salt solution was added in an amount sufficient to react with 25% of the silver nitrate. Then a 1.0% solution of sodium iodide was added in an amount sufficient to react with another 25% of the silver nitrate to produce a combination of silver chloride and silver iodide colloids from 50% of the silver nitrate.

Example 12

The process of Example 1 was repeated, except that

10 the NaCl salt solution was added in an amount sufficient to react with 25% of the silver nitrate to produce the poorly soluble silver chloride colloid. Then a 1.0% solution of sodium propionate was added in an amount sufficient to react with another 25% of the silver nitrate to produce the slightly water soluble silver

15 propionate colloid. Next, a 1.0% solution of sodium acetate was added in an amount sufficient to react with another 25% of the silver nitrate to produce the somewhat water soluble silver acetate colloid in combination with the poorly soluble silver chloride colloid and the slightly soluble silver propionate colloid

20 from 75% of the silver nitrate.

Example 13

The process of Example 12 was repeated, except that an additional amount of zinc iodide was added to convert 10% of

the remaining silver nitrate to a colloid of silver iodide. This produced a coating containing 15% silver nitrate, 25% of the somewhat soluble silver acetate colloid, 25% of the slightly soluble sodium propionate colloid, 25% of the poorly soluble silver chloride colloid, and 10% of the very poorly soluble silver iodide colloid, along with the soluble sodium nitrate and zinc nitrate salt products.

As shown by the above examples, any combination of additional salts in any combination of different amounts can be 10 used to convert some or all of the soluble oligodynamic metal salts into insoluble colloidal salts within a polymer composition.

#### Example 14

Somewhat water soluble silver salts, such as silver lactate or silver acetate, can be used alone or in combination with 15 the very soluble silver nitrate to produce other compounds that can have antiseptic activity. For example, silver acetate at a 4:1 molar ratio with zinc chloride produces 50% silver chloride colloid and the zinc acetate counter salt, which is also an antiseptic, and leaves 50% unreactive silver acetate. Similarly, other silver salts 20 can be used alone or in combination to produce multiple counter salts that have antiseptic or other desirable activity.

For example, the process of Example 2 was repeated except that a soluble combination of silver nitrate, silver acetate, and silver lactate was used in place of the 10% silver nitrate

solution. When the zinc chloride is added, a colloid of silver chloride is formed in the polymer composition and the soluble counter salts zinc nitrate, zinc acetate, and zinc lactate are produced. The zinc acetate and zinc lactate provide antiseptic activity in addition to the antimicrobial activity of the silver salts. In this example any metal salt other than zinc chloride which produces counter salts with the nitrate, acetate, and lactate that have a desired effect, such as antiseptic or antimicrobial activity, can be used. An example of such a salt is copper chloride.

Different oligodynamic salts have different water solubilities. This allows for tailoring of the composition to provide a specific release profile of the antimicrobial agent(s) from the composition. For example, sodium chloride, zinc iodide, sodium citrate, sodium acetate, and sodium lactate can be added to a coating composition containing silver nitrate to produce a coating which contains the water soluble salts silver nitrate and zinc nitrate, the somewhat water soluble salts silver lactate (67 mg/ml water) and silver acetate (10 mg/ml water), the slightly soluble salt silver citrate (0.3 mg/ml water), the poorly soluble salt silver chloride (0.002 mg/ml water), and the very poorly soluble salt silver iodide (0.00003 mg/ml water). By adjusting the proportions of salts having different solubilities in the composition, the release rate of the active oligodynamic agent(s) can be altered to provide a shorter or longer release profile over time.

For example, the process of Example 1 was repeated, except that in addition to the NaCl salt solution, 1% solutions of zinc iodide, sodium citrate, sodium acetate and sodium lactate were added, each in an amount sufficient to react with 15% of the

5 silver nitrate, to produce colloids of silver chloride, silver iodide, silver citrate, silver acetate, and silver lactate in the final coating composition, along with 25% unreacted silver nitrate, and the silver nitrate and zinc nitrate salt products. The difference in solubility of the different silver salts will produce different and

10 prolonged rates of silver ion release in the coating when exposed to body fluid.

Example 15

To form the coating composition for PVC catheters, a 3.3% solution of a polyether polyurethane-urea block copolymer

15 manufactured by Tyndale Plains-Hunter Ltd. (TPH) was prepared in THF. A 3.3% solution of Polyvinyl chloride (PVC) was then prepared in methylene chloride. The two solutions were then combined in equal amounts to provide a 50/50 ratio by weight of the two polymers in solution. A sufficient quantity of

20 10% silver nitrate ( $\text{AgNO}_3$ ) solution in alcohol was then added to the TPH/PVC polymer solution to produce a final silver concentration of approximately 5%, based on coating solids in the solution. A 1% zinc chloride solution in 75/25 mixture by weight of ethanol/water was added to the coating solution in an amount

sufficient to react with 50% of the  $\text{AgNO}_3$  to produce a colloid of the poorly water soluble salt  $\text{AgCl}$  from half of the  $\text{AgNO}_3$ . The  $\text{ZnCl}_2$  solution was added slowly to the polymer solution with stirring, and the solution began to turn cloudy with the formation 5 of the fine colloidal  $\text{AgCl}$ . The amount of water in the final coating solution was slightly less than about 1% of the total solvent weight. A PVC endotracheal tube was then coated by dipping it into the coating composition, followed by drying using standard methods. The finished coating contained both the 10 water soluble, and therefore fast releasing,  $\text{AgNO}_3$  and the poorly water soluble, and therefore slow releasing,  $\text{AgCl}$ .

Finally, it will be understood that the preferred embodiments have been disclosed by way of example, and that other modifications may occur to those skilled in the art without 15 departing from the scope and spirit of the appended claims.

## I Claim

1. An article of manufacture comprising a substrate and an antimicrobial polymer coating wherein the polymer coating comprises one or more polymers and a colloid wherein the colloid comprises salts of one or more oligodynamic metals  
5
2. The article of Claim 1 wherein the substrate is a medical device.
3. The article of Claim 2 wherein the medical device comprises a catheter, an endotracheal tube, a tracheostomy tube, a wound drainage device, a wound dressing, a stent, an implant, an intravenous catheter, a medical adhesive, a suture, a shunt, a glove, a condom, a contact lens, a gastrostomy tube, medical tubing, cardiovascular products, heart valves, pacemaker leads, a guidewire, or urine  
10 collection devices.
4. The article of Claim 3 wherein the medical device is a catheter.
5. The article of Claim 1 wherein the polymer comprises one or  
15 more hydrophilic polymers that are soluble in water or in an organic solvent containing water, one or more a hydrophobic polymers, or a combination of both hydrophilic and hydrophobic polymers.  
20

6. The article of Claim 5 wherein the hydrophilic polymer is selected from the group consisting of polyurethanes, polyvinylpyrrolidones, polyvinyl alcohols, polyethylene glycols, polypropylene glycols, polyoxyethylenes, polyacrylic acid, polyacrylamide, carboxymethyl cellulose, dextrans, polysaccharides, starches, guar, xantham and other gums, collagen, gelatins, biological polymers, and mixtures and copolymers thereof.  
5
7. The article of Claim 5 wherein the hydrophobic polymer is selected from the group consisting of polytetrafluoroethylene, polyvinyl chloride, polyvinylacetate, poly(ethylene terephthalate), silicone, polyesters, polyamides, polyureas, styrene-block copolymers, polymethyl methacrylate, polyacrylates, acrylic-butadiene-styrene copolymers, polyethylene, polystyrene, polypropylene, natural and synthetic rubbers, acrylonitrile rubber, cellulose, and mixtures, derivatives, and copolymers thereof.  
10
8. The article of Claim 1 wherein the colloid comprises at least one silver salt.  
20
9. The article of Claim 8 wherein the silver salt is selected from the group selected from the group consisting silver chloride, silver iodide, silver citrate, silver lactate, silver acetate, silver propionate, silver salicylate, silver bromide, silver ascorbate,

silver laurel sulfate, silver phosphate, silver sulfate, silver oxide, silver benzoate, silver carbonate, silver sulfadiazine, and silver gluconate.

10. The article of Claim 1 wherein the colloid comprises the salt  
5 of more than one oligodynamic metal.
11. The article of Claim 10 wherein the oligodynamic metal salts comprise salts having different solubilities in water.
12. The article of Claim 11 wherein the oligodynamic metal salts comprise silver lactate, silver acetate, silver citrate, silver  
10 chloride, and silver iodide.
13. An article of manufacture comprising one or more polymers and a colloid wherein the colloid comprises salts of one or more oligodynamic metals.
14. A composition comprising a polymer and a colloid wherein  
15 the colloid comprises salts of one or more oligodynamic metals.
15. The composition of Claim 14 wherein the colloid comprises at least one silver salt.
16. The composition of Claim 15 wherein the silver salt is selected from the group selected from the group consisting  
20 silver chloride, silver iodide, silver citrate, silver lactate, silver acetate, silver propionate, silver salicylate, silver

bromide, silver ascorbate, silver laurel sulfate, silver phosphate, silver sulfate, silver oxide, silver benzoate, silver carbonate, silver sulfadiazine, and silver gluconate.

17. The composition of Claim 14 wherein the colloid comprises  
5 the salt of more than one oligodynamic metal.
18. The composition of Claim 17 wherein the oligodynamic metal salts comprise salts having different solubilities in water.
19. The composition of Claim 18 wherein the oligodynamic metal salts comprise silver lactate, silver acetate, silver citrate,  
10 silver chloride, and silver iodide.
20. The composition of Claim 14 wherein the polymer comprises one or more hydrophilic polymers that are soluble in water or in an organic solvent containing water, one or more hydrophobic polymers, or a combination of both hydrophilic  
15 and hydrophobic polymers.
21. The composition of Claim 20 wherein the hydrophilic polymer is selected from the group consisting of polyurethanes, polyvinylpyrrolidones, polyvinyl alcohols, polyethylene glycols, polypropylene glycols, polyoxyethylenes, polyacrylic acid, polyacrylamide, carboxymethyl cellulose, dextrans, polysaccharides, starches, guar, xantham and other gums, collagen, gelatins, biological polymers, and mixtures and copolymers thereof.  
20

22. The composition of Claim 20 wherein the hydrophobic polymer is selected from the group consisting of polytetrafluoroethylene, polyvinyl chloride, polyvinylacetate, poly(ethylene terephthalate), silicone, polyesters, polyamides, polyureas, styrene-block copolymers, polymethyl methacrylate, polyacrylates, acrylic-butadiene-styrene copolymers, polyethylene, polystyrene, polypropylene, natural and synthetic rubbers, acrylonitrile rubber, cellulose, and mixtures, derivatives, and copolymers thereof.

10

23. A process for the manufacture of an antimicrobial composition comprising the steps of:

- 15 a) forming a solution, dispersion, or combination thereof of one or more polymers;
- b) forming in the polymer solution a colloid comprising contacting

  - 1) a solution of salt A which comprises one or more salts of oligodynamic metals, and
  - 2) a solution of salt B which comprises additional salts that reacts with salt A to form a fine colloidal precipitate.

20

24. The process of Claim 23 wherein the colloid in step (b) is formed separately and then combined with the polymer solution.

25. The process of Claim 23 wherein the colloid is formed *in situ* in the polymer solution.
26. The process of Claim 23 wherein salt A comprises salts of metals selected from the group consisting of silver, platinum, gold, zinc, copper, cerium, gallium, palladium, rhodium, iridium, ruthenium, and osmium.  
5
27. The process of Claim 23 wherein salt B comprises anions selected from the group consisting of acetates, ascorbates, benzoates, bitartrates, bromides, carbonates, chlorides, citrates, folates, gluconates, iodates, iodides, lactates, laurates, oxalates, oxides, palmitates, perborates, phenosulfonates, phosphates, propionates, salicylates, stearates, succinates, sulfadiazines, sulfates, sulfides, sulfonates, tartrates, thiocyanates, thioglycolates, and thiosulfates.  
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28. The process of Claim 23 wherein salt B comprises cations selected from the group consisting of calcium, sodium, lithium, aluminum, magnesium, potassium, and manganese.  
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29. The process of Claim 23 wherein the colloid formed in step (b) comprises at least one silver salt.  
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30. The process of Claim 29 wherein the silver salt is selected from the group consisting of silver chloride, silver iodide, silver citrate, silver lactate, silver acetate, silver propionate,

silver salicylate, silver bromide, silver ascorbate, silver laurel sulfate, silver phosphate, silver sulfate, silver oxide, silver benzoate, silver carbonate, silver sulfadiazine, and silver gluconate.

5 31. The process of Claim 23 wherein the colloid formed in step  
(b) comprises salts having different solubilities in water.

32. The process of Claim 31 wherein the colloid formed in step  
(b) comprises silver lactate, silver acetate, silver citrate, silver chloride, and silver iodide.

10 33. The process of Claim 23 wherein the polymer is a hydrophilic polymer, one or more a hydrophobic polymers, or a combination of both hydrophilic and hydrophobic polymers.

15 34. The process of Claim 23 wherein the polymer solution is formed by dissolving the polymer supercritical liquids.

35. A method for the manufacture of an article comprising dipping or spraying a substrate with a coating composition wherein the coating composition comprises polymers, monomers, or any combination thereof, and a colloid comprising salts of one or more oligodynamic metals.

20 36. A method for the manufacture of an article comprising the steps of

- 1) forming a solution, dispersion, of combination thereof of more or more polymers containing a colloid wherein the colloid comprises salts of one or more oligodynamic metals; and

5        2) drying the solution to create a solid polymeric article.

37. A method for the manufacture of an article comprising the steps of

- 1) forming a composition that comprises a solution, dispersion, of combination thereof of one or more polymers containing a colloid wherein the colloid comprises salts of one or more oligodynamic metals;

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- 2) drying the composition; and
- 3) processing the composition with the application of heat to form the article.

15 38. A method for the manufacture of an article comprising the steps of

- 1) forming a composition that comprises solution, dispersion, of combination thereof of one or more polymers containing a colloid wherein the colloid comprises salts of one or more oligodynamic metals;

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- 2) compounding the composition formed in step (1) with one or more polymers; and

- 3) processing the composition formed in step (2) with the application of heat to form the article.
- 5 39. A method for the manufacture of an article comprising dipping a form in a composition comprising polymers, monomers, or any combination thereof, and a colloid comprising salts of one or more oligodynamic metals.
- 10 40. A method for the manufacture of an article comprising casting a composition comprising polymers, monomers, or any combination thereof, and a colloid comprising salts of one or more oligodynamic metals.

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
21 June 2001 (21.06.2001)

PCT

(10) International Publication Number  
WO 01/043788 A3

(51) International Patent Classification<sup>7</sup>: A61L 27/34. (81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.

(21) International Application Number: PCT/US00/42372

(22) International Filing Date:  
29 November 2000 (29.11.2000)

(25) Filing Language: English (84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

(26) Publication Language: English

(30) Priority Data:  
09/461,846 15 December 1999 (15.12.1999) US

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Published:  
— with international search report

(88) Date of publication of the international search report:  
10 October 2002

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

A3

(54) Title: POLYMER COMPOSITIONS CONTAINING COLLOIDS OF SILVER SALTS

(57) Abstract: The present invention relates to antimicrobial compositions, methods for the production of these compositions, and use of these compositions with medical devices, such as catheters, and implants. The compositions of the present invention advantageously provide varying release kinetics for the active ions in the compositions due to the different water solubilities of the ions, allowing antimicrobial release profiles to be tailored for a given application and providing for sustained antimicrobial activity over time. More particularly, the invention relates to polymer compositions containing colloids comprised of salts of one or more oligodynamic metal, such as silver. The process of the invention includes mixing a solution of one or more oligodynamic metal salts with a polymer solution or dispersion and precipitating a colloid of the salts by addition of other salts to the solution which react with some or all of the first metal salts. The compositions can be incorporated into articles or can be employed as a coating on articles

WO 01/043788

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 00/42372

A. CLASSIFICATION OF SUBJECT MATTER	A61L27/34	A61L27/54	A61L29/08	A61L29/16	A61L31/10
IPC 7	A61L31/16				

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61L C23C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 320 908 A (SODERVALL BILLY V ET AL) 14 June 1994 (1994-06-14) claims; examples -----	1-40
X	DE 30 26 258 A (WAGSCHAL GMBH & CO KG FEINMECH) 28 January 1982 (1982-01-28) claims -----	1-40
A	WO 97 31709 A (UNIV NEW YORK) 4 September 1997 (1997-09-04) claims -----	1-40

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Date of the actual completion of the International search

2 November 2001

Date of mailing of the International search report

09/11/2001

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## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No  
PCT/US 00/42372

Patent document cited in search report		Publication date		Patent family member(s)		Publication date
US 5320908	A	14-06-1994	US	5395651 A	07-03-1995	
			US	5965204 A	12-10-1999	
			US	6224983 B1	01-05-2001	
			AT	134719 T	15-03-1996	
			AU	642872 B2	04-11-1993	
			AU	5473490 A	08-11-1990	
			CA	2016081 A1	04-11-1990	
			DE	69025495 D1	04-04-1996	
			DE	69025495 T2	11-07-1996	
			DK	400349 T3	18-03-1996	
			EP	0400349 A1	05-12-1990	
			EP	0693576 A1	24-01-1996	
			ES	2086328 T3	01-07-1996	
			FI	95816 B	15-12-1995	
			GR	3019899 T3	31-08-1996	
			JP	2645431 B2	25-08-1997	
			JP	3219085 A	26-09-1991	
			JP	3030687 B2	10-04-2000	
			JP	8325743 A	10-12-1996	
			JP	8333687 A	17-12-1996	
			NO	304746 B1	08-02-1999	
			US	5747178 A	05-05-1998	
DE 3026258	A	28-01-1982	DE	3026258 A1	28-01-1982	
WO 9731709	A	04-09-1997	AU	2061297 A	16-09-1997	
			EP	0883438 A1	16-12-1998	
			WO	9731709 A1	04-09-1997	
			US	6288076 B1	11-09-2001	
			US	6015816 A	18-01-2000	

